Remove

Figures Appear

Pages,Columns,Lines where

Relevant Passages or Relevant

Under the Paperwork Reduction Act of 1995, no persons are requ to a collection of information unless it contains a valid OMB control number.

INFORMATION DISCLOSURE	Application Number		10010200		
	Filing Date		2004-03-31		
	First Named Inventor DOU		JGLAS HOLBERG		
STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Art Unit		2816		
Not for Submission under 37 Of R 1.337	Examiner Name	HIEP	IEP NGUYEN		
	Attorney Docket Numb	er	CYGL-26,655		

U.S. PATENTS

Name of Patentee or Applicant

of cited Document

Kind

Code<sup>1</sup> Issue Date

	1	4429285	A	1984-01	-31	BRADSHAW					
	2	5159341	A	1992-10	-27	MCCARTNEY	ET AL.				
	3	6549066	B1	2003-04	-15	MARTIN					
If you wis	h to a	dd additional U.S. Pater	t citatio	n inform	ation pl	ease click the	Add button.		Add		
			U.S.P	ATENT	APPLIC	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>	Publica Date	tion	Name of Patentee or Applicant		Releva		Lines wher ges or Rele	
	1	20020190788	A1								
	ļ'	20020190788	AI.	2002-12	-19	RAMESH ET	AL.				
If you wis	_	dd additional U.S. Publi			-			d button	Add		
If you wis	_		shed Ap	plication	citation		please click the Adi	d buttor	Add		
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication FOREIC	citation	ent DOCUM	please click the Adi	e or	Remove Pages,Co where Rel	lumns,Line evant or Relevar	T=

Examiner Cite

Initial\* No Patent Number

# INFORMATION DISCLOSURE STATEMENT BY APPLICANT ( Not for submission under 37 CFR 1.99)

oplication Number		10816266			
ling Date		2004-03-31			
rst Named Inventor DOUG		GLAS HOLBERG			
t Unit		2816			
caminer Name HIEP		NGUYEN			
torney Docket Number		CYGL-26.655			

	2	0889597	₽	A2	1999-01-07	SHU				
If you wis	h to a	dd additional Foreign F	Patent Document	citation	information pl	lease click the Add butto	n Add			
			NON-PATE	NT LITE	RATURE DO	CUMENTS	Remove			
Examiner Initials*							Τs			
	1 SUPPLEMENTARY EUROPEAN SEARCH REPORT, EUROPEAN PATENT OFFICE, EP APPLICATION NO. 05 73 5518, 7/2/2007, pgs. 1-2.									
If you wish to add additional non-patent literature document citation information please click the Add button Add										
EXAMINER SIGNATURE										
Examiner	Signa	ture				Date Considered				
*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a										

F

E

See Kind Codes of USPTO Petent Documents at invest, ISPTO, DDL/or KMPEP 601.04. 2 Enter office that issued the document, by the involved code (WIPO) Standard S13.3 - For Lapraence parted rounders, the addication of the perior for the Emperor may precede the serial review of the cpatent document. If which office parted counters, and the comment of the perior for the perior may precede the serial review of the perior that office is a discussed on the document under WIPO Standard S11.6 if possible, "Applicant is to place a check mark here if Empirits frauques precision in statistics."

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

## INFORMATION DISCLOSURE STATEMENT BY APPLICANT ( Not for submission under 37 GFR 1.99)

Application Number		10816266			
Filing Date		2004-03-31			
First Named Inventor	DOUG	GLAS HOLBERG			
Art Unit		2816			
Examiner Name	HIEP	NGUYEN			
Attorney Docket Number		CYGL-26,655			

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s);

	That each item of information contained in the information disclosure statement was first cited in any communication
X	from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification
after making reasonable inquiry, no item of information contained in the information disclosure statement was known to
any individual designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure
statement. See 37 CFR 1.97(e)(2).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

RRIAN D WALKER

□ None

Name/Print

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Registration Number

37751

form of the signature.								
Signature	/BRIAN D. WALKER REG. # 37751/	Date (YYYY-MM-DD)	2007-08-14					

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 122 and 3T CFR 1.14. This collection is estimated to take it hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case: Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Peterta and Trademark Office, U.S. Operatment of Commence, P. 0. Dax 1436, Alexandris, V.S. 2231-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1459, Alexandria, V.S. 2231-1450.

### Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these records.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record perfains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
  application pursuant to 35 U.S.C. 12(2) to rissuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
  disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
  which became abandoned or in which the proceedings were terminated and which application is referenced by either a
  published application, an application open to public inspections or as issued patent.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.